

EXHIBIT C

1
2 UNITED STATES DISTRICT COURT
3 SOUTHERN DISTRICT OF NEW YORK

4 -----X
5 FEDERAL TRADE COMMISSION
6 and THE PEOPLE OF THE STATE
7 OF NEW YORK, by LETITIA
8 JAMES, Attorney General of
9 the State of New York,

10 Plaintiffs,
11 vs.

Case Number
1:17-cv-00124-LLS

12 QUINCY BIOSCIENCE HOLDING
13 COMPANY, INC., a
14 corporation, et al.,

15 Defendants.
16 -----X

17 VIDEOTAPED DEPOSITION OF PETER A. MALASPINA, Ph.D.

18 Taken Remotely

19 Friday, October 1, 2021
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22

23 Reported by

24 JEFFREY BENZ, CRR, RMR

25 JOB NO. 200428

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October 1, 2021

10:02 a.m.

Videotaped Deposition of PETER A. MALASPINA,
Ph.D., taken remotely, before Jeffrey Benz, a
Certified Realtime Reporter, Registered Merit
Reporter and Notary Public of the State of New
York.

A P P E A R A N C E S:

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A P P E A R A N C E S: (Ctd.)

ALSO PRESENT:

LARRY MOSKOWITZ, Videographer

JIMMY ROYER, Analysis Group

DAVID OVADIA, Federal Trade Commission

1 Malaspina

2 Where did you go to college and what
3 was -- what degree did you get?

4 A. I went to undergrad at Vassar. And
5 there I had a double major in math and
6 economics.

7 Q. Okay. And as part of your math or
8 economics degrees, did you study statistics at
9 all?

10 A. I did. Yes.

11 Q. Do you remember what you studied in
12 particular? That was a while ago.

13 A. It was. But I remember it. It was
14 one my favorite classes.

15 So, I think there was a -- like a
16 mathematical probability class which had a
17 sprinkling of statistics.

18 Then in economics, I took at least one
19 semester of econometrics, which is, you know, a
20 lot of statistics.

21 I took an advanced -- it's supposed to
22 be graduate-level mathematical statistics course
23 through the math department. And in terms of
24 pure stats classes, I think that's about it.

25 Q. Okay. And you mentioned econometrics.

Malaspina

Can you just explain what that is.

A. Yeah. So I'd just say it's a -- a branch of statistics where -- that invites more economic theory and economic concepts in. In a lot of cases I think that the term statistics and econometrics can be used interchangeably. Some people might think of econometrics as being -- having -- including more sophisticated economic modeling.

Q. Okay. And then after college what did you do?

A. For statistics?

Q. Just in life. Jobs, I guess, employment.

A. Yeah. I mean I took a year off. I drove across the country. But then, after that year, I went to get my Ph.D. at the University of North Carolina at Chapel Hill.

Q. What did you study there?

A. I studied economics.

Q. And as part of your studies in graduate school, did any of that involve statistics?

A. Yes.

1 Malaspina

2 Q. What kind of statistics did you study
3 there?

4 A. A lot.

5 So their first course was an advanced
6 probability and statistics course. I think it
7 was my first semester there.

8 Then -- I'm trying to remember the
9 sequence here because I actually took it out of
10 sequence slightly.

11 I think the second course, I ended up
12 take a third-year statistics course, which is
13 like an advanced financial econometrics. And it
14 was -- I took it out of sequence because they
15 only offered it every other year.

16 And, then, next I took an advanced
17 econometric theory course, which dealt with,
18 like, proofs, essentially mathematical proofs
19 using linear algebra of various statistical
20 models, econometric models.

21 And then I think the last course I
22 took was the advanced microeconomic models,
23 which would teach you a lot of applied --
24 basically walked through a bunch of applied
25 modeling and said, like, if the situation looks

1 Malaspina

2 like this, here's a model you might consider.

3 Yeah. So I think that's a -- it was
4 four courses in graduate econometrics.

5 Q. As part of either your undergraduate
6 or your graduate studies, did you use SAS
7 software or statistical software?

8 A. SAS. I don't think I used SAS. If I
9 did, it would be like a one off. Most of the
10 work in grad school was either done in MATLAB,
11 Stata, Excel, and there was Fortran. Ooph.

12 Q. I remember Fortran. But you don't
13 remember specifically using S-A-S or SAS?

14 A. No. I think my exposure to SAS came
15 in the consulting work that I did afterwards.

16 Q. Okay.

17 So after you graduated, got your
18 Ph.D., what did you do after that?

19 A. So, I went to work at Freeman,
20 Sullivan & Company, which they do, I call it,
21 electricity consulting out in San Francisco.

22 Q. And then what type of work did you do
23 for them?

24 A. So, I would call it just basically
25 economic consulting.

Malaspina

So let's see. I started off basically gathering data and running econometric analyses under the direction of other experts there.

So, I was learning about -- I would say mostly about electricity demand and sort of like daily demand, consumer behavior about demand.

And as I got more experience there, I started, you know, either conducting the analyses myself or having analysts under my direction conduct them, coming up to them with analysis plans, things like that.

Q. And I take it that your work for this company, Freeman, Sullivan, did not involve any statistical analysis of drug trials or supplement studies?

A. It did not.

Q. And did you do any statistical analysis of either drug trials or supplement clinical trials or supplement studies in college or graduate school?

A. Not that I can recall.

Yeah. Not that I can recall. I think there was, like, one time around the time I was

1 Malaspina

2 whereas what I was doing before was more like, I
3 would say, working for regulatory agencies,
4 slash, utilities. I had the opportunity to work
5 in, say, intellectual property and litigation.

6 And so it seemed like a cool
7 opportunity. My dissertation dealt with
8 intellectual property.

9 So, came out to Boston.

10 Q. And is that the Quantitative Economic
11 Solutions company?

12 A. Yes.

13 Q. And you started as a senior economist
14 there?

15 A. Yes.

16 Q. And what type of work did you do for
17 that company?

18 A. So, started off kind of learning the
19 ropes. I think playing a similar role to -- as
20 I -- how I did starting out at Freeman and
21 Sullivan, conducting analyses under the
22 direction of the expert, and then over time it
23 became more about, instead of carrying out the
24 direction of the expert, it was sort of like,
25 okay, here's the question. What do we do? You

Malaspina

figure it out.

I was developing econometric models, telling the analysts how to carry them out, writing a ton, writing up sections of reports.

Yeah, so that's a brief summary.

Q. What areas of focus did you have while you were working there?

A. Areas of focus. Like -- so intellectual property and antitrust I would say were two that stuck out.

Q. Okay. Anything else that you remember?

A. I mean, as a general topic, if you say damages calculation, I think there were some that weren't related to either intellectual property or antitrust. Like there were contract disputes, stuff like that.

Q. Did you work on any false advertising-type litigation?

A. Not that I can recall. Not at -- not at QES.

Q. Did any of your work there involve analysis of either drug trials or supplement studies?

Malaspina

A. I -- so it -- there -- a lot of the work we do is in pharmaceuticals, so there would be -- I think we would see reports that dealt with the clinical trial analyses. I don't ever remember, like, working with -- with clinical trial data itself.

Q. So in those pharmaceutical cases you were dealing with damages, for example, or antitrust issues?

MR. GLENNON: Objection. Form.

A. I --

Q. I guess what were you working on in those pharmaceutical cases?

A. Yeah. I -- I'll -- got to be careful not to give away confidential information, I guess --

Q. Sure.

A. -- but -- so a lot of things. But generally it related to intellectual property -- infringement and damages.

In some cases it would be like -- you know, like a commercial success report is where you're trying to show that a -- this is my understanding of the law.

1 Malaspina

2 So this is not my wheelhouse, but
3 you're trying to show that a product is
4 commercially successful, and that, according to
5 my understanding of the law, is an indicator
6 that it -- a patent is valid.

7 So, I think that's another -- in terms
8 of the pharmaceutical work we did, part of --
9 yeah, I think that's -- that's another area of
10 the pharmaceutical work we did.

11 And then there would be some contract
12 disputes in there also related to damages,
13 though.

14 Q. Okay. So you weren't studying or
15 analyzing the results of clinical trial data to
16 determine whether there were treatment effects
17 for the various pharmaceuticals or anything like
18 that?

19 A. No. The closest thing I would -- I
20 did something like that would be looking at
21 performance of a product over time and then
22 seeing whether or not like a -- you know what a
23 line extension is?

24 Q. No.

25 A. So when there's a change to a drug,

1 Malaspina

2 maybe they added an extended release formulation
3 of it.

4 Q. Sure.

5 A. You would see like, okay, does that
6 have an impact in demand or sales or number of
7 prescriptions.

8 So you might use an econometric model
9 to try to figure out that impact. But that's
10 not -- yeah, you -- I think you asked about
11 specifically clinical trial data. That's not
12 that.

13 Q. Right.

14 Okay. So then -- so -- right. So you
15 had -- you didn't have any involvement or work
16 on clinical trial data analyzing such studies?

17 A. No. I may have come across them in --
18 when we were, like, going through the documents
19 in some of these cases, but I myself did not
20 analyze clinical trial data there.

21 Q. And it's -- are you still working at
22 this company?

23 A. I -- I -- I am. I -- I haven't been
24 the entire time.

25 Q. Okay. So, I guess, what happened

1 Malaspina

2 after -- you were a senior economist for how
3 long?

4 A. I forget really. I want to say three
5 years?

6 Q. Okay.

7 A. Maybe four.

8 Q. What did you do after that?

9 A. So I was senior economist. At some
10 point I became a VP. And then I went from there
11 to New York to work at the New York Attorney
12 General's office as their chief economist.

13 And the reason for the confusion is --
14 and then after two years there I came back to
15 QES. So --

16 Q. I see. So you left Quantitative
17 Economic Solutions to go to the New York
18 Attorney General's office?

19 A. Yes.

20 Q. And what does the chief economist for
21 the New York AG do?

22 A. What don't they do?

23 A lot of things. I mean it's one of
24 the -- one of the cool parts about that job is
25 you sort of make it what you want it to be.

1 Malaspina

2 I'm not sure if that's how it was
3 intended, but that was my experience there.

4 I think primarily you're there to
5 offer guidance to the attorney general, and any
6 of the attorneys working for her or him. And
7 that can take a variety of forms.

8 Sometimes I was doing a calculation
9 quickly to get a rough idea of what -- you know,
10 what might be at stake for a certain case that
11 the office was thinking about.

12 And other times I would be sitting in
13 the room with lawyers negotiating a potential
14 settlement. Sometimes I was helping the data
15 analytics team actually program analyses,
16 working with Excel.

17 It was -- yeah, there was just so many
18 facets to the job. I could probably go on for a
19 long time, but I don't know if that's what
20 you're looking for.

21 Q. You said you worked with Excel. Did
22 you work with SAS while you were at the New
23 York -- with the New York AG?

24 A. Yeah. I think there was one thing I
25 did in SAS.

1 Malaspina

2 They had -- there -- the way they were
3 set up this, they had access to all the -- every
4 program you could ever possibly want.

5 Q. You remember working with it one time?

6 A. I think so, yeah.

7 I think it was a situation where an
8 opposing expert had used it, and so it was --
9 make -- I think -- you know, when you first look
10 at something you use the program that they're --
11 that they're using.

12 Q. And is that something you generally
13 try to do, use the same program that opposing
14 experts use?

15 A. When you're first looking at it. You
16 know, you want to see what did they see when
17 they were programming their model.

18 Q. And what types of litigations did you
19 work on at the AG's office?

20 MS. MATUSCHAK: I just want to caution
21 the witness not to reveal anything that
22 might be privileged, work product or
23 confidential.

24 Q. Sure.

25 You can answer.

1 Malaspina

2 A. A lot of -- do you mean like the types
3 of --

4 Q. I guess areas of, like, antitrust or
5 other types of --

6 A. Yeah. So a lot of it was in
7 antitrust, but since I was the chief for the
8 whole office, I got dragged into environmental.
9 There were, I'd say, a finance -- personal care
10 matters.

11 Q. Sorry. I didn't hear that last.

12 A. Health care.

13 Q. Health care. Yeah.

14 A. Yeah. There was one matter where I
15 provided an expert report. It was a health
16 care, slash -- I think there was a bureau of tax
17 or taxation. I forget what it is.

18 But, yeah, there was -- I was kind of
19 involved in a bunch of different areas of all
20 the office's work. I think the only thing that
21 I -- well, only broad bureau that I didn't get
22 involved with -- and this may actually be a
23 division, not a bureau, was like the criminal
24 division.

25 Q. You did have any involvement with

Malaspina

know, the names of everybody involved in this matter. I'm just trying to be -- it would be very surprising to me if I was involved in anything like that.

Q. Okay. Anything involving Quincy or this litigation?

A. Yes.

Q. All right. And what did you do after your time at the New York Attorney General's office?

A. I came back to QES.

Q. Is that where you still are today?

A. Yes.

Q. What types -- did you do any different type of work when you went back to QES than you did -- than what we discussed earlier?

A. I mean, subject matter is mostly the same. There's a lot of antitrust and intellectual property.

There's been some -- so some of the work, I guess, that would be unique this time around. I've done some consulting where I've been helping pharmaceutical companies develop their -- I would say like a licensing plan and

1 Malaspina

2 for licensing some of their intellectual
3 property with other firms.

4 And that's not in the context of
5 litigation. It's more just saying, here are
6 some things you -- you know, think about in the
7 long run for your -- for how you want to, you
8 know, structure a royalty rate or a loss of
9 exclusivity term, something like that.

10 Q. Okay. And since you've been back at
11 QES, have you been involved in the analysis of
12 any clinical trial data for either drugs or
13 supplements?

14 A. Again, if I -- I may have seen
15 documents that dealt with clinical trials, but
16 I -- I wasn't analyzing the underlying clinical
17 trial data.

18 Q. Okay. I think I forgot to ask
19 earlier, but either your first time at QES or
20 since you've been back have you used SAS?

21 A. Yes.

22 Q. In what capacity?

23 A. A lot. I think the first thing I did
24 when I got to QES, the first big thing, was --
25 I'm just trying to be careful I don't give away

Malaspina

confidential information. But we were working with very large health -- health insurance claims data set. And I quickly realized that my -- my tool of choice, Stata, wasn't going to do it. There was a memory issue where you can't get enough information into it.

So I -- you know, my -- I think SAS might be my -- one of my second tools of choice.

So I said, okay, well, let's figure out whether SAS can do it. And that was when I learned that -- at least at that time, SAS is a big advantage in terms of dealing with large data sets, the way it handles memory. So, we used SAS extensively on that health insurance matter, which went on for a couple years.

And there -- I think at that point, just as -- it's a relatively small firm, right? So if somebody like me or another seniorish manager type feels strongly about a methodology, we kind of -- it becomes the rule of how to handle things.

So I think we sort of developed a best practice of when we've got large data sets, at least the initial phases of importing data,

1 Malaspina

2 merging it, any of the big steps where you're
3 trying to maybe even chop up the data again or
4 append large data sets, that's all done in SAS.

5 And, in fact, I got to -- I guess, by
6 necessity, I could build, like, the hardware
7 or -- I didn't build it physically myself, but
8 got to, you know, choose all the attributes of a
9 SAS server that we had built so we could handle
10 large data sets like that. Again.

11 It was specifically for SAS.

12 Q. Right.

13 Would you say that you used SAS while
14 you were at QES mostly for data manipulation as
15 opposed to statistical modeling?

16 A. It's hard to say most. You mean,
17 like, time spent programming?

18 Q. Sure. Like your -- I mean your use of
19 SAS. Would you describe it as mostly to
20 manipulate the data or to do statistical
21 modeling?

22 A. Yeah. I'm not trying to split hairs,
23 but when you do the data manipulation on large
24 data sets, sometimes it's like -- I can program
25 the import and merging and then export in five

1 Malaspina

2 the record. The time is 11:05 a.m.

3 (Recess from 11:05 a.m. to

4 11:13 a.m..)

5 THE VIDEOGRAPHER: The time is

6 11:13 a.m. We are back on the record.

7 Q. Just to go back for a second,

8 Dr. Malaspina, to your work history and

9 education, you said that you did use SAS in your

10 work with your current company, Quantitative

11 Economic Solutions; is that correct?

12 A. Yes.

13 Q. In using SAS at that company, did you

14 use the proc mixed statement at all? Or

15 procedure?

16 A. Where -- I'm not sure if I've used it

17 before the present matter. I've used several

18 things like it. It's possible that I used it at

19 some point during that. It's hard to remember

20 every single instance where I've used SAS.

21 Q. You don't recall, sitting here today,

22 using it prior to this litigation?

23 A. Again, you know, there's -- when it

24 comes to econometrics and statistics, there are

25 many different programs and many different tools

1 Malaspina

2 A. I do.

3 Q. And is it fair to say that this is
4 your main criticism of Dr. Beales' model?

5 Does this kind of sum it up?

6 A. I wouldn't say that. Yeah, I -- I
7 think there are several criticisms. I'm not
8 sure this captures everything.

9 Q. Okay. But one of your criticisms is
10 that, in your view, Dr. Beales' model assumes no
11 correlation in error terms across the
12 observations, right?

13 A. Yes.

14 Q. And just to clarify, it's your opinion
15 that the model should allow for correlation and
16 test scores for the same individual; is that
17 right?

18 A. So, when it comes to my opinions about
19 the model, it's really about what did Dr. Beales
20 say he was trying to do, and how did he describe
21 what he did. And so, I'm just trying to
22 demonstrate that what he actually did, in his
23 analysis that he presented, wasn't that.

24 So, I'm not saying that this is the
25 right or wrong way to conduct an analysis of the

Malaspina

underlying data, merely correcting what the --
serious mistakes made by Dr. Beales.

Q. I get it. But one of those mistakes
that you're saying he made, in your opinion that
he made, is his model assumes no correlation in
error terms across any observations, right?

A. Yes. He said that he was going to
address correlation, and he didn't. So that's a
mistake.

Q. And the correlation he wanted to
address would be within individual results. In
other words, if somebody does well on one test,
you might expect them to do well on another
test?

A. That was one form of the correlation
that he was trying to address.

Q. Right. So that would be within
subject or within participant correlation?

A. That would be one form, yes.

Q. Take a look at Figure 2 of your
report, which is on the top of page 10. And
this is a snapshot of Dr. Beales' SAS code; is
that right?

A. Yes.

1 Malaspina

2 to what went into your report and why, then
3 I would ask me -- instruct you not to
4 answer.

5 A. Yeah, I don't think -- I think this is
6 a, you know, determination I made by myself.

7 It didn't go in my report, because it
8 wasn't necessary to reach the opinions I was
9 making.

10 Q. Okay. Do you remember what the
11 outcome was when you ran the model with the
12 unstructured variance structure for the R
13 matrix?

14 A. I want to be careful here, because
15 it's been a bit. And these are obviously fine
16 details. I think it was on structuring the R.
17 It may have been a broader unstructuring of what
18 is the -- the -- essentially the -- the V
19 matrix, which is the combination of R and Z.

20 But my recollection was, is that
21 the -- the program didn't even -- it couldn't
22 find a solution. It basically hung up, which is
23 not -- it's not too surprising when you're --
24 when you're using a very flexible model and
25 you're giving a lot of freedom, which -- that's

Malaspina

sort of what you're trying to do when you're --
freeing up choice of variances.

Sometimes the model can get confused.
And again, it's probably also a symptom of mixed
model and SAS being very flexible. So, maybe
not that surprising.

Q. When you say "hung up," is that sort
of a technical issue, not enough sort of
computing power to finish, or what?

A. I mean -- technical issue. I mean it
could be one of those things where just the --
the way SAS tries to reach solution, it gets
caught in the local -- a local minimum, stuck at
the bottom of a well, basically, and when it's
trying to find a solution and it can't, it can't
go -- there's nowhere else to go from there.

So I'm not sure. It's just a question
of like computational firepower. Like if I put
it on a NEC supercomputer it might solve.

Does that answer the question?

Q. Sure. Did you -- did you try any
other types of variance structures for the R
matrix other than going all the way to an
unstructured matrix?

1 Malaspina

2 A. I don't think so.

3 Q. For example, did you try a compound
4 symmetry?

5 A. I don't think so.

6 I mean, let me just make sure I
7 remember what compound symmetry looks like.

8 Q. Sure.

9 A. Yeah. I'm fairly certain I never
10 attempted compound symmetry.

11 Q. Okay. Why don't we take another five
12 minutes, and then we'll continue.

13 A. Great.

14 THE VIDEOGRAPHER: We are going off
15 the record. The time is 12:07 p.m.

16 (Recess from 12:07 to 12:16.)

17 THE VIDEOGRAPHER: We are back on the
18 record. The time is 12:16 p.m.

19 Q. Great. So before the break we were
20 talking about another analysis that you did or
21 tried to do using the unstructured variance R
22 matrix.

23 Were there any other -- and sorry.

24 And that analysis did not find its way
25 into your report. Are there any other analyses

1 Malaspina

2 that you performed or attempted to perform that
3 are not in your report?

4 A. Yes. So there's probably other -- as
5 you're programming this stuff up and trying to
6 make sure that your, you know, results are
7 correct, you may run small permutations of stuff
8 just to see what's going on with the data.
9 Stuff like that.

10 Things that weren't -- obviously
11 weren't essential to my opinion, things like
12 looking at the data in summary form. So like,
13 actually literally just pulling all the data
14 into -- I may have started in Excel, but just
15 like looking at a pivot table and what does the
16 stuff even look like.

17 Those are important steps for just
18 getting yourself comfortable and knowing what
19 you're working with. But again, it's -- this is
20 sort of preliminary and -- and not essential for
21 the ultimate opinion.

22 RQ MR. SUDEN: Okay. I think we're
23 going to ask that you produce all analyses
24 that you performed in working on -- on
25 whether it's included in your report or

1 Malaspina

2 more time?

3 Q. Do you agree that -- do you agree that
4 the only reason your modified model finds no
5 statistical significance for the joint tests is
6 because she used a completely different
7 approach -- i.e., the bootstrap method -- to
8 model the variance, covariants metrics?

9 MR. GLENNON: Objection.

10 A. So, the reason why -- my corrections
11 to Dr. Beales produces different results is
12 because it's -- doing what he said he was doing
13 and what he failed to program, so that has to do
14 in part with the way you model -- you allow the
15 model to account for correlation across
16 equations and also across -- within individuals.

17 Q. Okay. I asked you -- and I guess,
18 just to be clear, Dr. Beales doesn't use or
19 mention the bootstrap model in his report,
20 right?

21 A. He talks about random effects, and
22 then does not use them, and so the bootstrap
23 method is one way to incorporate the random
24 effects.

25 Q. Right. But he doesn't do that or

1 Malaspina

2 recollection. But that's sort of where I'm
3 sitting here. I think it's highly unlikely I
4 received anything related to Quincy.

5 Q. Okay.

6 MR. SUDEN: I'm going to that ask the
7 New York AG's office produce any documents
8 that were sent to, cc'd, bcc'd,
9 Dr. Malaspina, while he was the chief
10 economist that have anything to do with
11 Quincy or the issues in this case.

12 MS. MATUSCHAK: I can just represent
13 to you on the record that Dr. Malaspina did
14 not work on Quincy matters while he was at
15 the AG.

16 MR. SUDEN: Right. I appreciate that,
17 but the doctor seems to be saying that he
18 may have received emails, on/off emails or
19 something that have to do with Quincy or
20 Prevagen. So we would just ask that you
21 confirm that either no such emails were
22 sent to him or, if they were, that he
23 produce them.

24 MS. MATUSCHAK: We can deal with that
25 off the record.

C E R T I F I C A T E

STATE OF NEW YORK)
) Ss.:
COUNTY OF NEW YORK)

I JEFFREY BENZ, a Certified Realtime Reporter, Registered Merit Reporter and Notary Public within and for the State of New York, do hereby certify:

That PETER A. MALASPINA, Ph.D., the witness whose examination is hereinbefore set forth, was duly sworn by me and that this transcript of such examination is a true record of the testimony given by such witness.

I further certify that I am not related to any of the parties to this action by blood or marriage; and that I am in no way interested in the outcome of this matter.

IN WITNESS WHEREOF, I have hereunto set my hand this 13th of October, 2021.



JEFFREY BENZ, CRR, RMR

**UNITED STATES DISTRICT COURT
SOUTHERN DISTRICT OF NEW YORK**

Case No. 1:17-cv-00124-LLS

FEDERAL TRADE COMMISSION and
THE PEOPLE OF THE STATE OF NEW
YORK, by LETITIA JAMES, Attorney
General of the State of New York,

Plaintiffs,

v.

QUINCY BIOSCIENCE HOLDING
COMPANY, INC., a corporation;

QUINCY BIOSCIENCE, LLC, a limited
liability company;

PREVAGEN, INC., a corporation
d/b/a/ SUGAR RIVER SUPPLEMENTS;

QUINCY BIOSCIENCE
MANUFACTURING, LLC, a limited
liability company; and

MARK UNDERWOOD, individually and as
an officer of QUINCY BIOSCIENCE
HOLDING COMPANY, INC., QUINCY
BIOSCIENCE, LLC, and PREVAGEN,
INC.,

Defendants.

I, Peter A. Malaspina, hereby make the following corrections to the transcript of my deposition, which occurred on October 1, 2021:

PAGE	LINE(S)	CORRECTION	REASON
56	18	“I test” to “ith test”	Typographical error
57	13	“01” to “0/1”	Typographical error
90	16	“on structuring” to “unstructuring”	Typographical error

